

Amendment to the Specification:

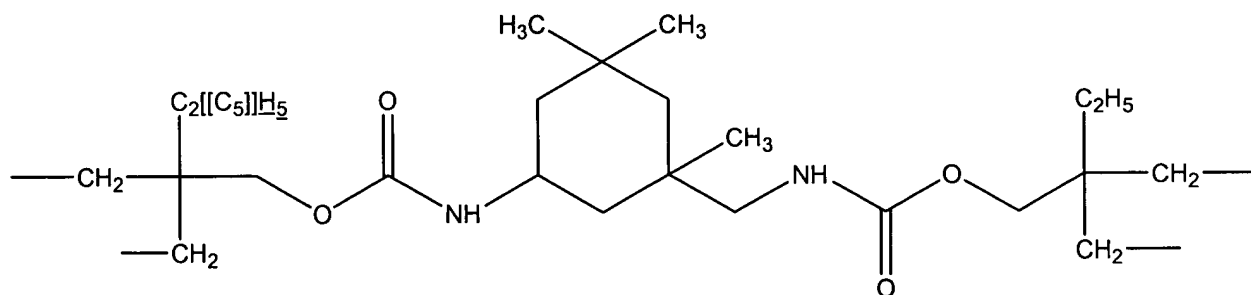
Please add the following new paragraph to the first line of the specification:

This application is a divisional of Application No. 10/031,983, filed January 25, 2002, which is the National Stage of International Application No. PCT/EP00/06931, filed July 20, 2000.

Please amend the specification at page 1 by replacing the last line with the following:

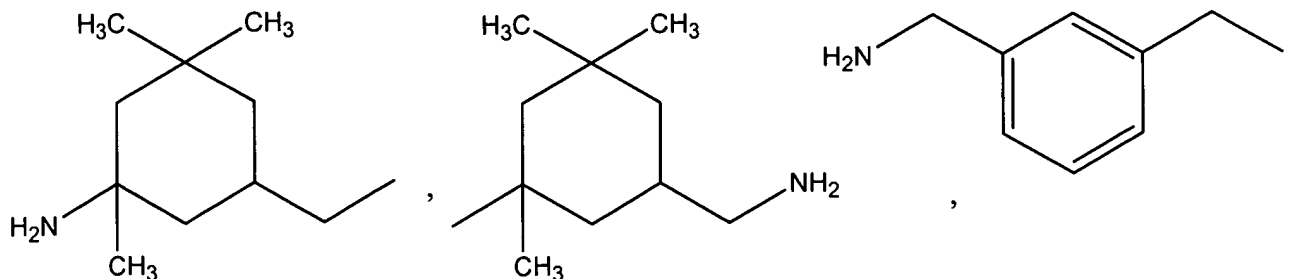
X is -O-, $[-COO-]$, $-C(=O)O$ or $-CHR_4-$, with R_4 and R_3 together forming an ethylene group,

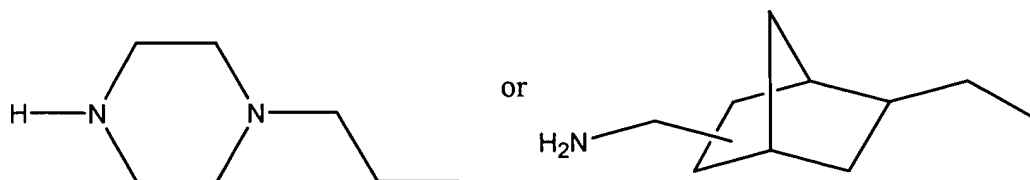
Please amend the specification at page 3 by replacing the formula for the tetravalent radical with the following formula:



Please replace the sixth paragraph on page 4 of the specification with the following paragraph:

Special preference is given to compounds of formulae Ia and Ib wherein $[[R_1]]$ R_5 is n-butyl, n-octyl, cyclohexyl, benzyl, 2-aminoethyl, 4-(aminomethyl)pentyl, 5-amino-2-methylpentyl, 3-dimethylaminopropyl, 3-methylaminopropyl, 4-aminocyclohexyl or a radical of formula $-CH_2CH_2NHCH_2CH_2NH_2$,





Please amend the specification at page 5 by replacing the last paragraph with the following:

The episulfide of formula IIIa is then dissolved in an aprotic or protic organic solvent and, under inert gas (argon or nitrogen), reacted with the amine $R_5\text{-NH-R}_2$, the amount of the amine $R_5\text{-NH-R}_2$ preferably being so selected that there are from 1 to 10 $[[\text{NH}]]$ amine groups for one episulfide group. Preferred solvents are alcohols (e.g. methanol, ethanol, tert-butanol) and aromatic hydrocarbons, for example toluene and xylene

Please amend the specification at page 7 by replacing the first full paragraph with the following:

Episulfides can, for example, also be ~~synthesised~~ synthesized from the corresponding epoxides by reaction with triphenylphosphine sulfide.

Please replace the first full paragraph on page 11 with the following paragraph:

As component $[[A]]$ C of the substance mixtures in accordance with the invention, preference is given to cycloaliphatic and aliphatic amines, especially the amines of formulae $R_5\text{-NH-R}_2$ and $E\text{-(NHR}_2\text{)}_{m+1}$ used for preparation of the polymercaptopolyamines according to the invention.

Please replace the paragraph bridging pages 12 and 13 with the following paragraph:

I. Preparation of compounds of formula $[[I]]$ Ia and Ib

a) General procedure for the preparation of polyepisulfides:

~~The polyepoxide of formula II is~~ epoxy compounds of formula IIa or IIb are dissolved in an amount of solvent that is from 0.5 to 5 times the

amount of the ~~polyepoxide~~ epoxy compound and is stirred, under nitrogen, with thiourea or alkali-metal or ammonium thiocyanate (0.8-1.2 equivalents of sulfur per epoxy equivalent) at 60-100°C until the epoxy content has fallen to nearly zero.

After separating off the by-products by means of filtration, extraction or phase separation, the polyepisulfide is isolated as a result of concentration by evaporating off the solvent.

b) General procedure for the preparation of polymercaptopolyamines:

The polyepisulfide is dissolved in an amount of solvent that is from 0.5 to 5 times the amount of the polyepisulfide and under nitrogen and with vigorous stirring, is combined with the amine (either R_5-NH-R_2 or $E-(NHR_2)_{m+1}$), which likewise has been dissolved in an amount of solvent that is from 0.5 to 5 times the amount of the amine. The amount of amine is selected so that there are from 1 to 10 $[[NH_2]]$ amine groups for one episulfide group. After stirring at 60-100°C for from 0.2 to 3 hours, the solvent is distilled off under reduced pressure. To isolate the polymercaptopolyamine of formula $[[I]]$ Ia or Ib, the excess amine reagent is removed by means of vacuum distillation at elevated temperature.

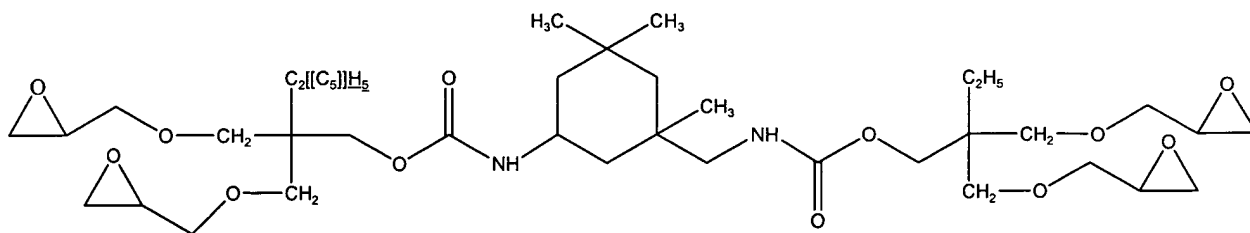
In one embodiment of the invention, the excess amine is not removed and the mixture of the excess amine $[[R_1-NH_2]]$ and the polymercaptopolyamine of formula $[[I]]$ Ia or Ib is used as a hardener for epoxy resins.

Please replace the paragraph bridging pages 13 and 14 with the following paragraph:

In accordance with the above-mentioned procedure, polymercaptopolyamines according to the invention (~~Examples I.1-I.19~~) (Examples I.1-I.46) are prepared from the following diamines R_1-NH_2 amines and epoxides of formula II epoxy compounds of formula IIa or IIb:

BA:	n-butylamine
OA:	n-octylamine
CYA:	cyclohexylamine

BZA:	benzylamine
MBA:	methylbutylamine
DMDP:	N,H-demethyl-1,3-diaminopropane
MDP:	N-methyl-1,3-diaminopropane
DACY:	1,2-diaminocyclohexane
AEP:	N-2-aminoethylpiperazine
DETA:	diethylenetriamine
IPD:	isophoronediamine
MXDA:	meta-xylylenediamine
DYTEK-A:	1,5-diamino-2-methylpentane
NBDA:	isomeric mixture of 2,5- and 2,6-bis(aminomethyl)norbornane
EDA:	ethylenediamine
epoxide 1:	liquid diglycidyl ether of bisphenol A having an epoxy content of 5.25-5.4 eq./kg
epoxide 2:	liquid mixture of diglycidyl ether of bisphenol A and diglycidyl ether of bisphenol F having an epoxy content of 5.5-5.8 eq./kg
epoxide 3:	1,4-bis(hydroxymethyl)cyclohexane diglycidyl ether
epoxide 4:	epoxy phenol novolak having an epoxy content of 5.6-5.8 eq./kg
epoxide 5:	diglycidyl ether of hydrogenated bisphenol A
epoxide 6:	tetraglycidyl ether of formula



epoxide 7:	di(β -methylglycidyl) ether of bisphenol A
epoxide 8:	trimethylolpropane triglycidyl ether

epoxide 9: hexahydrophthalic acid diglycidyl ester (epoxy content:
5.6-6.2 eq./kg)
epoxide 10: phenyl glycidyl ether

Please replace the first footnote on page 16 with the following footnote:

^{*)} mixture of amine $[[R_1-NH_2]]$ and polymercaptopolyamine; excess $[[R_1-NH_2]]$ amine
not distilled off

Please amend the Abstract by replacing the tenth line with the following:

X is -O-, $[-COO-]$ $-C(=O)O$ or $-CHR_4-$, with R_4 and R_3 together forming an ethylene
group,